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**Discovery and characterization of a sulfoquinovose mutarotase using kinetic analysis at equilibrium by exchange spectroscopy**

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16   **Abstract**

17   Bacterial sulfoglycolytic pathways catabolise sulfoquinovose (SQ), or glycosides thereof, to  
18   generate a three-carbon metabolite for primary cellular metabolism and a three-carbon  
19   sulfonate that is expelled from the cell. Sulfoglycolytic operons encoding an Embden-  
20   Meyerhof-Parnas (EMP)-like or Entner-Doudoroff (ED)-like pathway harbour an  
21   uncharacterized gene (*yihR* in *Escherichia coli*; *PpSQ1\_00415* in *Pseudomonas putida*) that is  
22   upregulated in the presence of SQ and has been annotated as an aldose-1-epimerase and which  
23   may encode an SQ mutarotase. Our sequence analyses and structural modelling confirmed that  
24   these proteins possess mutarotase-like active sites with conserved catalytic residues. We over-  
25   expressed the homologue from the sulfo-ED operon of *Herbaspirillum seropedicaea* (*HsSQM*)  
26   and used it to demonstrate SQ mutarotase activity for the first time. This was accomplished  
27   using NMR exchange spectroscopy (EXSY), a method that allows chemical exchange of  
28   magnetization between the two SQ anomers at equilibrium. *HsSQM* also catalyzed the  
29   mutarotation of various aldohexoses with an equatorial 2-hydroxy group, including D-  
30   galactose, D-glucose, D-glucose-6-phosphate, and D-glucuronic acid, but not D-mannose.  
31   *HsSQM* displayed only 5-fold selectivity in terms of efficiency ( $k_{\text{cat}}/K_{\text{M}}$ ) for SQ versus the  
32   glycolysis intermediate glucose-6-phosphate (Glc-6-P), however its proficiency [ $k_{\text{uncat}} /$   
33   ( $k_{\text{cat}}/K_{\text{M}}$ )] for SQ was 17,000-fold better than for Glc-6-P, revealing that *HsSQM* preferentially  
34   stabilises the SQ transition state.

## 35 **Introduction**

36 Various prokaryotes metabolise the sugar sulfoquinovose (SQ) to sulfolactaldehyde (SLA) and  
37 dihydroxyacetone phosphate (DHAP), via an Embden-Meyerhof-Parnas (EMP)-like pathway  
38 [1], or pyruvate, via an Entner-Doudoroff (ED)-like pathway [2, 3] (Figure 1). While the DHAP  
39 or pyruvate feed into primary metabolic pathways, SLA is converted to 2,3-  
40 dihydroxypropanesulfonate (DHPS) or sulfolactate (SL) by the sulfo-EMP and sulfo-ED  
41 pathways, respectively, and excreted from the cell. Yet sulfoquinovose is rarely encountered  
42 as a free sugar in nature; rather it is liberated from the plant sulfolipid  $\alpha$ -sulfoquinovosyl  
43 diacylglycerol (SQDG), or its delipidated form  $\alpha$ -sulfoquinovosyl glycerol (SQGro), by the  
44 action of glycoside hydrolases termed sulfoquinovosidases (SQases) [4]. SQases are retaining  
45 glycosidases, and result in the initial formation of  $\alpha$ -SQ, which can undergo mutarotation to  $\beta$ -  
46 SQ at an unknown rate. The anomeric preferences of the immediate downstream enzymes that  
47 utilize SQ (SQ isomerase for the sulfo-EMP pathway; SQ dehydrogenase for the sulfo-ED  
48 pathway) are unknown.

49 All proteins comprising a sulfo-ED or sulfo-EMP pathway are typically encoded within  
50 a single gene cluster. These clusters usually include an SQase, which highlights that SQ  
51 glycosides are important natural feedstocks for sulfoglycolytic pathways [1, 2]. The gene  
52 clusters also encode a conserved uncharacterized protein, annotated as an aldose-1-epimerase,  
53 which likely catalyses SQ mutarotation: an enzyme activity yet to be reported. Mutarotases are  
54 widely distributed enzymes that facilitate the rapid mutarotation of aldoses to enhance flux  
55 through metabolic pathways when enzymes acting on reducing sugars are specific for a single  
56 anomer [5, 6].

57 A classical approach to studying mutarotases involves measuring rates of mutarotation  
58 by polarimetry [7-10]. These assays are limited by an inability to prepare pure samples of a  
59 single anomer of many reducing sugars. Indeed, to date it has not been possible to obtain SQ  
60 as a single anomer, which makes this approach of limited use for studying putative SQ  
61 mutarotases (SQMs). Alternative approaches to studying mutarotases have employ coupled  
62 assays in which only one anomer acts as substrate for a secondary enzyme [11, 12] or a  
63 chemical reaction (e.g. bromine oxidation [13]), or use a glycosidase to prepare a single anomer  
64 in solution prior to addition of the mutarotase [12]. The high rate of uncatalyzed mutarotation  
65 under standard conditions often renders these assays technically demanding, and a requirement  
66 for different coupling enzymes for each substrate undermines their generality. An alternative  
67 approach is to study reaction rates at equilibrium. NMR spectroscopy is ideally suited to this

68 approach using the technique of exchange spectroscopy (EXSY) [14, 15]. EXSY involves  
69 chemical shift labelling of the spin population of nuclei at one site within a substrate, followed  
70 by a chemical reaction that changes the chemical environment of individual nuclei resulting in  
71 magnetization transfer to the new site, and finally sampling of the magnetization states of the  
72 nuclei in the substrate and product. Because of the spectral resolution of NMR spectroscopy  
73 and the potential to conduct two-dimensional variants, EXSY can be used to study  
74 unidirectional reactions at equilibrium. Several reports have described the development of  
75 saturation transfer and inversion transfer NMR methods for analysis of equilibrium exchange  
76 rates for mutarotases [16-19].

77         Here, we disclose the first measurement of SQ mutarotase activity, using an enzyme  
78 from *Herbaspirillum seropedicaea* (*HsSQM*) and 2D EXSY, and analyze its selectivity for  
79 various reducing sugars with and without an anionic substituent at C6. *HsSQM* exhibited a  
80 broad spectrum of activity for sugars with an equatorial hydroxyl group at C2 and with, or  
81 without, charge at C6. 1D EXSY was used to measure reaction rates to obtain Michaelis-  
82 Menten kinetics for *HsSQM* with SQ and glucose-6-phosphate, a common cytoplasmic  
83 metabolite, revealing an approximate 5-fold preference for SQ as a substrate. Sequence and  
84 structural analyses revealed *HsSQM* belongs to the galactose mutarotase-like Structural  
85 Classification Of Proteins (SCOP) with strictly conserved active site residues that are proposed  
86 to be involved in substrate binding and catalysis.

87

## 88 **Materials and Methods**

### 89 **Reagents**

90 D-Glucose (Glc), D-mannose (Man), D-galactose (Gal), D-glucuronic acid (GlcA) and D-  
91 glucose-6-phosphate (Glc-6-P) were purchased from Sigma Aldrich. D-Sulfoquinovose was  
92 purchased from MCAT GmbH (Germany, <http://www.MCAT.de>). 4-Nitrophenyl  $\alpha$ -D-  
93 sulfoquinovoside and sulfoquinovosidase have been described previously [4].

### 95 **Cloning, expression and purification of HsSQM**

96 A gene encoding the *Herbaspirillum seropedicaea* strain AU14040 protein WP\_069374721.1  
97 was codon harmonised for *E. coli*, synthesised and cloned into the pET29 vector using the *NdeI*  
98 and *XhoI* restriction sites to provide pET29-HsSQM (see Supporting Information). This  
99 plasmid was transformed into chemically competent 'NEB Express' *E. coli*, plated onto LB-  
100 agar (50  $\mu$ g/ml kanamycin) and incubated at 37 °C for 16 h. A single colony was used to  
101 inoculate 10 ml of LB media containing 50  $\mu$ g/ml kanamycin followed by incubation at 37 °C  
102 for 16 h. This culture was used to inoculate 1000 ml of "S-broth" (35 g tryptone, 20 g yeast  
103 extract, 5 g NaCl, pH 7.4) containing 50  $\mu$ g/ml kanamycin, which was incubated with shaking  
104 (250 rpm) at 37°C until it reached an OD<sub>600</sub> of 0.7. The culture was cooled to room temperature,  
105 IPTG added to a final concentration of 100  $\mu$ M, and then incubated with shaking (200 rpm) at  
106 18 °C for 19 h. The cells were harvested by centrifugation at 17,000 g for 20 min at 4 °C then  
107 resuspended in 40 ml of binding buffer (50 mM NaPi, 500 mM NaCl, 5 mM imidazole, pH 7.5)  
108 containing protease inhibitor (Roche complete EDTA-free protease inhibitor cocktail) and  
109 lysozyme (0.1 mg/ml) by nutating at 4 °C for 30 min. Benzonase (1  $\mu$ l) was added to the  
110 mixture and lysis was effected by sonication. The lysate clarified by centrifugation (17,000 g  
111 for 20 min at 4 °C), the supernatant filtered (0.45  $\mu$ m) and loaded onto a 1 ml HiTrap TALON  
112 column (GE Healthcare). The column was washed with 15 ml binding buffer and the protein  
113 was eluted using elution buffer (50 mM NaPi, 500 mM NaCl, 500 mM imidazole, pH 7.5).  
114 Fractions containing the protein of interest (as determined by SDS-PAGE) were further  
115 purified by size exclusion chromatography on a HiLoad 16/600 Superdex 75 column using 50  
116 mM NaPi, 150 mM NaCl, pH 7.5. Fractions containing the protein of interest were further  
117 purified on a MonoQ 5/50 column with protein bound in buffer A (20 mM Tris, pH 7.5) and  
118 eluted with a linear gradient from 100% buffer A to 100% buffer B (20 mM Tris, NaCl 500  
119 mM, pH 7.5) over 15 ml at 1 ml/min. Protein yield was  $\approx$ 5 mg per litre of culture.



## NMR magnetization-transfer experiments

The theoretical background of NMR for the study of kinetics of chemical exchange has been well documented [14, 20] and only a brief overview is presented. The interconversion of anomers by mutarotation is described by a simple two-state equilibrium:



For a species exchanging between isomers, the dependence upon time ( $t$ ) of the longitudinal nuclear magnetizations of corresponding nuclei from the  $\alpha$  and  $\beta$  anomer spin populations ( $I_\alpha$ ) and ( $I_\beta$ ), corresponding to the integrals, are given by the Bloch-McConnell equation:

$$\frac{d}{dt} \begin{pmatrix} I_\alpha \\ I_\beta \end{pmatrix} = \begin{pmatrix} -R_\alpha - k_1 & k_{-1} \\ k_1 & -R_\beta - k_{-1} \end{pmatrix} \begin{pmatrix} I_\alpha \\ I_\beta \end{pmatrix} + \begin{pmatrix} R_\alpha & 0 \\ 0 & R_\beta \end{pmatrix} \begin{pmatrix} I_\alpha^0(t) \\ I_\beta^0(t) \end{pmatrix} \quad (2)$$

Where  $R_\alpha$  and  $R_\beta$  are the longitudinal relaxation rates, characterizing the return of the magnetizations towards their respective equilibrium values,  $I_\alpha^0$  and  $I_\beta^0$ ;  $k_1$  and  $k_{-1}$  are the extrinsic rate constants for the forward and reverse reactions for each anomer,  $\alpha$  and  $\beta$  respectively. The solution of this equation gives the intensity of the transferred magnetization at  $\tau_{\text{mix}}$ :

$$\begin{pmatrix} I_\alpha(\tau_{\text{mix}}) \\ I_\beta(\tau_{\text{mix}}) \end{pmatrix} = \exp \left[ \begin{pmatrix} -R_\alpha - k_1 & k_{-1} \\ k_1 & -R_\beta - k_{-1} \end{pmatrix} \tau_{\text{mix}} \right] \times \begin{pmatrix} I_\alpha(0) - I_\alpha^0 \\ I_\beta(0) - I_\beta^0 \end{pmatrix} + \begin{pmatrix} I_\alpha^0 \\ I_\beta^0 \end{pmatrix} \quad (3)$$

Under the condition of chemical equilibrium, the net flux is zero. However, the unidirectional fluxes remain and are equal in both directions. Thus,

$$k_1[\alpha]^{\text{eq}} = k_{-1}[\beta]^{\text{eq}} \quad (4)$$

And

$$K_{\text{eq}} = \frac{[\beta]^{\text{eq}}}{[\alpha]^{\text{eq}}} = \frac{k_1}{k_{-1}} = \exp\left(-\frac{\Delta G}{RT}\right) \quad (5)$$

To obtain the rate constants,  $k_1$  and  $k_{-1}$ , at a given concentration ratio of the substrate and enzyme, a series of 1D  $^1\text{H}$  EXSY spectra with mixing time  $\tau_{\text{mix}}$ , ranging from 5 ms to 1.5 s were acquired using a 1D selective NOESY pulse sequence. For each substrate concentration, the normalized integrals of the substrate ( $\text{H1}\alpha$ ) and product ( $\text{H1}\beta$ ) anomeric signals are plotted against mixing time, providing buildup and decay curves, respectively, as described previously [21]. The data were then fitted to a second-order polynomial, an approximation to Eq 3, which is valid for short mixing times. The tangent at  $\tau_{\text{mix}} = 0$ , the so-called initial rate approach [22], provides estimates of both exchange-rate constants  $k_1$  and  $k_{-1}$ .

### Michaelis–Menten kinetics

The equilibrium-exchange kinetics (reaction rate,  $v^{\text{eq}}$ ) of a mutarotase-mediated reaction can be expressed as:

$$v^{\text{eq}}([\alpha]) = \frac{V_{\text{max}}^{\text{eq}}[\alpha]}{K_{\text{M}}^{\text{eq}} + [\alpha]} \quad (9)$$

$V_{\text{max}}^{\text{eq}}$  is the maximum rate of reaction, reached under saturating equilibrium exchange conditions when  $[\alpha] \gg K_{\text{M}}^{\text{eq}}$ . The rate of the mutarotation process can be expressed in terms of the Michaelis-Menten equation:

$$v^{\text{eq}} = k_1[\alpha]^{\text{eq}} = \frac{V_{\text{max}}^{\text{eq}}[\alpha]}{K_{\text{M}}^{\text{eq}} + [\alpha]} \quad (10)$$

$k_{\text{cat}}$  is calculated by dividing  $V_{\text{max}}^{\text{eq}}$  by the enzyme concentration.

### 2D $^1\text{H}$ - $^1\text{H}$ EXSY experiments

2D  $^1\text{H}$ - $^1\text{H}$  EXSY spectra for all substrates (SQ, Glc-6-P, Glc, Gal, Man, GlcA) were acquired at 25 °C on a Bruker Avance II 800 spectrometer equipped with a TXI cryoprobe using the standard 2D NOESY/EXSY pulse sequence (noesyphpr). Spectra were collected with 4 and 8 scans in the absence and presence of enzyme, respectively. A mixing time,  $\tau_{\text{mix}}$  of 1.1 s was used for all the experiments. Spectra were processed and analysed using TOPSPIN (version 3.2 Bruker) and  $^1\text{H}$  chemical shifts were referenced indirectly to DSS at 0 ppm via the  $\text{H}_2\text{O}$  resonance, 4.77 ppm at 25 °C. Samples were prepared in  $\text{D}_2\text{O}$  buffer consisting of 50 mM



sodium phosphate, 150 mM NaCl (pD 7.5) with a substrate concentration 5 mM in the absence or presence of 1.51  $\mu$ M HsSQM. pD was measured using a pH meter and the relationship  $pD = pH + 0.4$ .

## 1D $^1H$ EXSY experiments

### *Michaelis-Menten parameters*

1D  $^1H$  EXSY spectroscopic studies for SQ and Glc-6-P were performed at 25 °C on a Bruker Avance III 600 spectrometer equipped with a TCI cryoprobe using a 1D selective NOESY pulse sequence (selnogg, Bruker). Samples were prepared in D<sub>2</sub>O buffer consisting of 50 mM sodium phosphate, 150 mM NaCl (pD 7.5) using 1.50  $\mu$ M HsSQM at substrate concentrations ranging from 0.5-15.0 mM (for SQ) or 0.5-30.0 mM (for Glc-6-P). pD values were calculated using the following relationship:  $pD = pH + 0.4$ . For each sample, 1D  $^1H$  EXSY spectra with mixing time,  $\tau_{mix}$ , ranging from 5 ms to 1.5 s were acquired with number of scans varying between 32 and 256 depending on the concentration of substrate. A recycle delay of 13.4 s (c.a. 3-5 times of measured  $^1H$   $T_1$  relaxation times) between scans was used for the acquisition of 1D  $^1H$  EXSY spectra. Spectra were subsequently processed and analysed using TOPSPIN (version 3.2 Bruker). Kinetic data for conversion of the  $\beta$ -anomer of SQ to the  $\alpha$ -anomer were obtained by selectively inverting the resonance of H5 $\beta$  at 3.72 ppm using a Gaussian-shaped pulse of 20 ms, and *vice versa*, for conversion of the  $\alpha$ -anomer to the  $\beta$ -anomer, the same selective pulse was applied to the resonance of H5 $\alpha$  at 4.15 ppm. Similarly, for the conversion of the  $\beta$ -anomer of Glc-6-P to the  $\alpha$ -anomer, kinetic data were obtained by selectively inverting the signal for H1 $\beta$  at 4.57 ppm. A build-up curve for the  $\beta$ -anomer could not be obtained since the signal for H1 $\beta$  at 4.57 ppm was affected by the nearby HOD peak at 4.77 ppm (see SI data for representative  $^1H$  NMR spectra). Instead, the rate constant for the conversion of the  $\alpha$ -anomer to the  $\beta$ -anomer were calculated using Eq 4. Rates for each concentration were calculated using the Prism 6 software package (GraphPad Scientific Software). Data were fitted to a second-order polynomial function as described previously by Aski *et al.* [21].

### *pD dependence of activity for HsSQM*

The Michaelis-Menten parameter  $k_{cat}/K_M$  was measured for Glc-6-P mutarotation in D<sub>2</sub>O buffer consisting of 50 mM citrate/phosphate, 150 mM NaCl at a range of pD values (5.6, 6.1, 6.5, 7.0, 7.5, 8.1, 9.0, 9.4, 9.8, 10.4 and 10.9) at 25 °C. Reactions were initiated by adding 1.48-2.96  $\mu$ M HsSQM to Glc-6-P (5.0 mM) in buffer and the rate measured by 1D  $^1H$  EXSY as

described above with mixing time,  $\tau_{\text{mix}}$ , ranging from 5 ms to 1.0 s and number of scans from 64 and 128. Kinetic data were obtained for the conversion of the  $\beta$ -anomer of Glc-6-P to the  $\alpha$ -anomer, by selectively inverting the signal for H1 $\beta$  at 4.57 ppm.  $k_{\text{cat}}/K_M$  and  $pK_a$  values were calculated using the Prism 6 software package (Graphpad Scientific Software). pH dependence was fit to the following function:

$$y = \frac{k_{\text{cat}}}{K_M} \left[ \left( \frac{1}{1 + \left( \frac{10^{-\text{pH}}}{10^{-\text{p}K_{a1}} + \frac{10^{-\text{p}K_{a2}}}{10^{-\text{pH}}}} \right)} \right) \right] + c \quad (11)$$

### **Uncatalyzed mutarotation rate measurement**

The uncatalyzed rate constant for SQ mutarotation was measured using polarimetry on a Jasco DIP-1000 digital polarimeter equipped with Na 589 nm lamp and 100.00 mm cell, using a sulfoquinovosidase to generate  $\alpha$ -SQ from 4-nitrophenyl  $\alpha$ -D-sulfoquinovoside (PNPSQ). Analysis by  $^1\text{H}$  NMR spectroscopy and thin layer chromatography revealed that hydrolysis of PNPSQ was complete within 5 min. SQase (final concentration 2.13  $\mu\text{M}$ ) was added to a solution of PNPSQ (final concentration 12.4 mM) in buffer in a final volume of 2 ml. Buffers consisted of: 50 mM sodium phosphate, 150 mM NaCl in  $\text{H}_2\text{O}$  (pH 7.1); 50 mM sodium phosphate, 150 mM NaCl in  $\text{D}_2\text{O}$  (pD 7.5); or 10-50 mM sodium phosphate, 2 M NaCl in  $\text{D}_2\text{O}$  (pD 7.5). The reaction mixture was transferred to the polarimetry cell and after 5 min the mutarotation rate was monitored continuously at  $26 \pm 1$   $^\circ\text{C}$  by reading specific rotation at various times. The rate constant was calculated using the Prism 6 software package (Graphpad Scientific Software). Data were fitted to a one phase decay function,  $t_{1/2} = \ln(2)/k$ .

For mutarotation as described in equation 1, the rate of change in the concentration of the  $\alpha$ -anomer has two contributions: it is depleted by the forward reaction at rate  $k_1[\alpha]$  and is replenished by the reverse reaction at rate  $k_{-1}[\beta]$ . The net rate of change is therefore:

$$\frac{d[\alpha]}{dt} = -k_1[\alpha] + k_{-1}[\beta] \quad (12)$$

The solution of this first-order differential equation is:

$$[\alpha] = \left\{ \frac{k_{-1} + k_1 e^{-(k_1 + k_{-1})t}}{k_1 + k_{-1}} \right\} [\alpha]_0 \quad (13)$$

And thus the first order decay constant ( $k$ ) is related to the forward and reverse rates as:

$$k = k_1 + k_{-1} \quad (14)$$

Using equation (5), which relates the equilibrium constant to the forward and reverse rates, this shows that:

$$K^{eq} = \frac{[\beta]^{eq}}{[\alpha]^{eq}} = \frac{k_1}{k-k_1} \quad (15)$$

## Results

Our initial efforts to characterize SQ mutarotases focused on expressing *YihR* from *E. coli* (which utilizes the sulfo-EMP pathway) [1], and *PpSQ1\_00415* from *P. putida* SQ1 (which utilizes the sulfo-ED pathway) [2] in an *E. coli* expression system. However, despite screening several expression constructs and conditions, only a poor yield of low quality protein was ever obtained. Thus, we turned our attention to other putative SQ mutarotases from various bacteria that possess sulfoglycolytic gene clusters and succeeded in obtaining WP\_069374721.1 (hereafter *HsSQM*) from *Herbaspirillum seropedicaea* in high yield and purity. *H. seropedicaea* is a nitrogen-fixing endophytic bacterium capable of colonizing the intercellular spaces of grasses such as rice and sugar cane [23], and contains a predicted sulfo-ED operon analogous to that of *P. putida* SQ1 but lacking in synteny (Figure 2A) [2]. A sequence alignment of these three putative sulfoquinovose mutarotases, as well as two structurally characterized hexose mutarotases, is provided in Figure 2B: *HsSQM* shares 37% similarity (19% identity) with *E. coli* *YihR*, and 50% similarity (35% identity) with *P. putida* SQ1 *PpSQ1\_00415* (SI Table 1). All three proteins retain the highly conserved residues of other hexose mutarotases (His92, His162 and Glu254 in *HsSQM*). The equivalent residue to *HsSQM* His162 in galactose mutarotase from *E. coli* has previously been proposed to be involved in substrate binding [24], whereas the equivalent residues to His92 and Glu254 are proposed to act in roles of general acid and general base, respectively, in the first half of the reaction leading to the acyclic aldehyde, in galactose mutarotases from both *L. lactis* and *E. coli* [24, 25].

The iTASSER server [26-28] provided a homology model of *HsSQM* with a C-score of 0.87, suggesting it to be a good approximation for the protein's native fold. This was compared to existing structures by using TM-align [29, 30] and returned only other (predicted) mutarotase structures. Greatest structural similarity was found between the *HsSQM* model and

a galactose mutarotase-like protein from *Clostridium acetobutylicum* (PDB 3OS7, Figure 3A), with a RMSD of 1.55 Å between structural models, despite both proteins sharing <15% identity. Structural alignment of the *HsSQM* homology model and the galactose mutarotase domain of gal10 from *S. cerevisiae* with D-galactose bound (PDB 1Z45) [31] reveals that the conserved active site residues His92, His162 and Glu254 in the homology model are positioned appropriately for catalysis (Figure 3B).

2D  $^1\text{H}$ - $^1\text{H}$  EXSY was used to assess whether *HsSQM* can catalyze mutarotation. A 2D pulse sequence equivalent to that used for a 2D  $^1\text{H}$ - $^1\text{H}$  NOESY experiment was employed to provide a visual representation of the chemical exchange network [15, 20]. A similar approach was utilized by Graille and coworkers to study a hexose-6-phosphate mutarotase [32], building on work by Balaban and Ferretti for the study of anomerization/isomerization of Glc-6-P by phosphoglucose isomerase using 2D  $^{31}\text{P}$ - $^{31}\text{P}$  EXSY [17], and earlier work by Kuchel and coworkers, who studied mutarotation catalyzed by porcine mutarotase using  $^{13}\text{C}$ - $^{13}\text{C}$  EXSY [16]. In this approach, all nuclei are excited using a  $90^\circ$  excitation pulse, which is allowed to evolve over a  $t_1$  period. A second  $90^\circ$  pulse is then applied to rotate the  $y$ -component of magnetization onto the  $z$ -axis, and a mixing interval  $\tau_{\text{mix}}$ , allows magnetization transfer through chemical exchange. A final  $90^\circ$  pulse creates transverse magnetization in the  $xy$  plane, which is detected. Molecules that undergo chemical interconversion display cross-peak signals between a signal from the substrate along one axis of the 2D spectrum, and the corresponding nucleus in the product on the other axis. Careful choice of mixing time to be shorter than that required for chemical exchange by spontaneous mutarotation can allow qualitative detection of enzyme-catalyzed mutarotation in a single NMR experiment. Figure 4A shows that a solution of SQ displays H1 of the  $\alpha$ - and  $\beta$ -anomers as independent sets of signals, and upon addition of *HsSQM* off-diagonal cross-peaks appear between the anomeric protons. This data provides evidence that chemical exchange is occurring and that *HsSQM* is catalyzing the mutarotation of SQ.

To qualitatively explore the substrate specificity of *HsSQM*, we assessed the ability of the enzyme to catalyze mutarotation of other simple hexoses (Figures 4B-F). *HsSQM* could catalyze the mutarotation of D-glucose-6-phosphate and D-glucuronic acid, showing that the sulfonate group is not required and that *HsSQM* can tolerate other anionic groups. A set of simple aldohexoses was also studied. *HsSQM* catalyzed mutarotation of D-glucose and D-galactose, showing that the enzyme is tolerant of either stereochemistry at C4. However, using this pulse sequence we could not detect *HsSQM*-mediated mutarotation of D-mannose on the

NMR time-scale, demonstrating the preference of *HsSQM* for substrates with an equatorial C2-hydroxyl group.

While these experiments provide insights into the substrate specificity of *HsSQM*, simple hexoses are not present at any appreciable concentration within bacterial cells, as shown for *E. coli* grown on various carbon sources [33]. Other than SQ, the only other physiologically plausible substrate for *HsSQM* is glucose-6-phosphate (Glc-6-P): an abundant primary metabolite produced by bacteria grown on glucose, glycerol or acetate [33]. To explore the selectivity of *HsSQM* for SQ and Glc-6-P, we used EXSY to determine kinetic parameters under conditions of equilibrium exchange. While it is possible to use 2D methods to determine rates, accurate determination requires *a priori* knowledge of optimum mixing times and a lengthy experiment, and so we elected to use the alternative 1D EXSY approach. Measurement of the equilibrium-exchange rate constants was achieved by selective inversion of a distinctive signal from one anomer and then monitoring the return to equilibrium intensity of signals from both anomers. For quantitative determination of rates, the selective irradiation must be of sufficient power and duration to invert the targeted magnetization, yet should not directly affect the equilibrium magnetization of the other spin population. Due to the chemical exchange of the anomers, the inverted magnetization redistributes into the un-irradiated site. The magnetization can then be sampled by normal methods. Figures 5A,B show build-up curves of H1 $\alpha$  arising from the irradiation of H1 $\beta$  of SQ and Glc-6-P, respectively; and Figures 5C,D show the respective decay curves from the same experiment for H1 $\beta$ . Also shown is the tangent at  $\tau_{\text{mix}} = 0$ , which provides the exchange-rate constant. Figures 5E,F show Michaelis-Menten plots for conversion of the  $\beta$ -anomers of SQ and Glc-6-P, respectively. While SQ exhibited saturation kinetics, allowing calculation of  $k_{\text{cat}}$ ,  $K_{\text{M}}$  and  $k_{\text{cat}}/K_{\text{M}}$  values, Glc-6-P did not and thus only  $k_{\text{cat}}/K_{\text{M}}$  could be calculated. As the equilibrium concentrations of the two anomers of each substrate are known, this data allows calculation of kinetic parameters for the reverse reaction. The complete set of kinetic parameters for the forward and reverse mutarotation reactions are shown in Table 1, and reveals that *HsSQM* displays an approximate 5-fold selectivity for SQ over Glc-6-P in terms of the  $k_{\text{cat}}/K_{\text{M}}$  values.

**Table 1.** Michaelis Menten kinetic parameters for *HsSQM* catalyzed mutarotation of SQ and Glc-6-P under conditions of equilibrium exchange.

substrate	$k_{\text{cat}}$ (s <sup>-1</sup> )	$K_{\text{M}}$ (mM)	$k_{\text{cat}}/K_{\text{M}}$ (M <sup>-1</sup> s <sup>-1</sup> )
$\alpha$ -SQ	$(3.08 \pm 0.19) \times 10^2$	$1.00 \pm 0.20$	$(3.07 \pm 0.44) \times 10^5$

$\beta$ -SQ	$(3.37 \pm 0.20) \times 10^2$	$2.08 \pm 0.34$	$(1.62 \pm 0.18) \times 10^5$
$\alpha$ -Glc-6-P	—	—	$(5.87 \pm 0.57) \times 10^4$
$\beta$ -Glc-6-P	—	—	$(3.26 \pm 0.31) \times 10^4$

To gain insight into the rate enhancement achieved by *HsSQM* we measured the rate of mutarotation of SQ by polarimetry using a coupled assay. Incubation of 4-nitrophenyl  $\alpha$ -D-sulfoquinovoside with a retaining sulfoquinovosidase occurs with retention of configuration to afford  $\alpha$ -SQ. Monitoring the rate of mutarotation of  $\alpha$ -SQ in 50 mM phosphate buffer revealed a half-life of 50 min (SI Figure 1). By comparison, mutarotation of  $\alpha$ -Glc-6-P has been reported to occur at 25 °C with a half-life of 6.0 s [34, 35]. However, phosphate buffer is known to catalyze mutarotation.[36] In order to measure the rate of mutarotation in phosphate-free conditions, and to subsequently allow comparison with enzyme-catalyzed rates measured in D<sub>2</sub>O we measured mutarotation rates of SQ in D<sub>2</sub>O at various concentrations of phosphate at pseudo-constant ionic strength achieved with 2 M NaCl (Figure 6A). Extrapolation of this data to [phosphate] = 0 mM, gave a predicted mutarotation rate ( $k$ ) of  $3.87 \times 10^{-5} \text{ s}^{-1}$  ( $0.00232 \text{ min}^{-1}$ ) corresponding to a half-life of 299 min. Using the equilibrium constant  $K^{\text{eq}} = 1.5$  allows calculation of forward and reverse rates of  $k_1 = 2.3 \times 10^{-5} \text{ s}^{-1}$  and  $k_{-1} = 1.5 \times 10^{-5} \text{ s}^{-1}$ . A similar calculation for Glc-6-P mutarotation ( $K^{\text{eq}} = 1.8$ ) using the published data gave forward and reverse rates of  $k_1 = 7.7 \times 10^{-2} \text{ s}^{-1}$  and  $k_{-1} = 4.3 \times 10^{-2} \text{ s}^{-1}$ ; strictly speaking these 'uncatalyzed' rates are for  $k_{(\text{H}_2\text{O})} + k_{(\text{H}^+)}[\text{H}_3\text{O}^+] + k_{(\text{HO}^-)}[\text{HO}^-]$ .

The above data allow us to calculate the catalytic proficiency of *HsSQM* for catalyzing the mutarotation of SQ and Glc-6-P. The proficiency constants [ $k_{\text{uncat}} / (k_{\text{cat}}/K_{\text{M}})$ ] for *HsSQM* are  $7.6 \times 10^{-11} \text{ M}$  for  $\alpha$ -SQ and  $1.3 \times 10^{-6} \text{ M}$  for  $\alpha$ -Glc-6-P, revealing that *HsSQM* binds the SQ transition state some 17,000-fold tighter than for Glc-6-P and is thus best considered a sulfoquinovose mutarotase. We note that the mutarotation rate for Glc-6-P was measured in H<sub>2</sub>O, whereas all other measurements were performed in D<sub>2</sub>O, thus a solvent isotope effect may somewhat confound this analysis; nonetheless the broad conclusions still apply.

The pD dependence of activity for *HsSQM* was determined using 1D <sup>1</sup>H EXSY for Glc-6-P. At a concentration of 5 mM, Glc-6-P does not show saturation (unlike for SQ), and so rates measured at this concentration provide an estimate of  $k_{\text{cat}}/K_{\text{M}}$ . Figure 6 shows the pD dependence of activity is bell-shaped, with a broad maximum of activity of pD 7-9. The simplest interpretation of these results is that it arises from ionization of two catalytically important residues with pK<sub>a</sub> values of  $6.3 \pm 0.2$  and  $10.3 \pm 0.2$ . The ionization of the acidic limb



presumably reflections ionization of the general acid Glu254, and the basic limb ionization of the general base His92. By comparison, the intrinsic  $pK_a$  values for the ionizable side-chains of Glu and His are 4.5 and 6.4, respectively [37]. In both cases, and particularly the latter, these intrinsic  $pK_a$  values are perturbed, presumably because of the active-site environment including charge-charge and charge-dipole interactions, as well as desolvation effects [37]. The pD dependence of this enzyme is similar to the pH dependence of green pepper mutarotase [34], but broader than that of *E. coli* galactose mutarotase [38].

## Discussion

Many enzymes involved in the metabolism of free sugars are stereospecific for one anomer of their substrate. For example, yeast galactokinase specifically phosphorylates  $\alpha$ -galactose to produce  $\alpha$ -D-galactose-1-phosphate [39], glucose dehydrogenase from *Bacillus megaterium* uses only  $\beta$ -D-glucose as a substrate [40]; yeast phosphoglucose isomerase only uses  $\alpha$ -D-glucose-6-phosphate as substrate [41]; and yeast phosphomannose isomerase is specific for  $\alpha$ -D-mannose-6-phosphate [42]. For these enzymes, access to the appropriate substrate anomer would be rate-limiting *in vivo* if the process relied on spontaneous mutarotation. Mutarotases ensure that there is rapid equilibration between anomers to eliminate this metabolic bottleneck. While some mutarotases have been linked to specific metabolic pathways [6], it has been difficult to assign many others to a single substrate or metabolic process [32]. It is unclear if the inherent promiscuity of many mutarotases confers an advantage to their hosts or simply impose no cost to fitness.

Sulfo-EMP and sulfo-ED gene clusters both encode putative mutarotases, homologous to HsSQM, that are upregulated in response to growth on SQ [1, 2], suggesting that their primary function is as an SQ mutarotase. We show here that HsSQM from the sulfo-ED cluster of *H. seropedicaea* can catalyze SQ mutarotation with a greater  $k_{cat}/K_M$  value than for the possible alternative substrate Glc-6-P, which is itself an important metabolite that feeds into the Embden-Meyerhof-Parnas and Entner-Doudoroff glycolytic pathways, as well as the pentose phosphate pathway. It is possible that the activity of the enzyme on Glc-6-P is advantageous to a bacterium transitioning between growth on SQ and growth on Glc. Furthermore, due to the broad substrate tolerance of many mutarotases, it is possible that other enzymes may play a role in catalysing SQ mutarotation within the cell, though any enzyme that does so is unlikely to be transcriptionally regulated by SQ concentration as HsSQM and its homologs are.

The substrate specificity of *HsSQM* bears similarities to the aldose-1-epimerase from *E. coli* (galM), which is tolerant to functional group changes at C6, and stereochemistry changes at C4 (Gal) but not at C2 (Man) [9]. On the other hand it is distinguished from yeast ymr099c, which encodes a hexose-6-phosphate mutarotase with activity on Glc-6-P, Gal-6-P and Man-6-P and thus tolerates stereochemical inversion at C2 [32]. Sequence alignments and structural modelling reveal that *HsSQM* likely acts through a mechanism conserved with all other mutarotases. The substrate tolerance of *HsSQM* for substituent variation at C6 lies in contrast to *E. coli* SQase, which exhibited negligible activity on  $\alpha$ -glucosides due to its specialized set of conserved residues that recognize the sulfonate group of SQ glycosides [4].

The effectiveness of an enzyme as a catalyst can be quantified by measuring the rate enhancement it provides relative to the rate of the uncatalyzed reaction [43]. This work reveals that uncatalyzed SQ mutarotation is a relatively slow reaction, with the rate for  $\alpha$ -SQ some 3400-fold lower than that of  $\alpha$ -Glc-6-P. On the other hand, *HsSQM* catalyzes mutarotation with a  $k_{\text{cat}}/K_{\text{M}}$  value around 5-fold higher than for Glc-6-P. Combining these values reveals that *HsSQM* is approximately 17,000-fold more proficient as a catalyst in catalyzing the mutarotation of SQ and Glc-6-P. As enzymes achieve their rate enhancement through selective stabilization of the transition state relative to the ground state, these data suggest that the affinity for the transition state of SQ mutarotation is approximately 17,000-fold greater than that for Glc-6-P mutarotation. As has been noted by others, the unusually high rate of mutarotation of Glc-6-P is much greater than that of other hexoses [44], and appears to be a result of neighboring group participation by the pendant phosphate group [36]; our data suggests that the sulfonate group of SQ does not provide neighbouring group participation.

Understanding the precise role of SQMs in the sulfo-EMP and sulfo-ED pathways will require knowledge of the substrate specificities for upstream and downstream processes. Upstream processes include SQ importers and SQases. For organisms grown on a mixture of SQ anomers, the SQ importers may exhibit a preference for only one SQ anomer; an SQM may be required for re-establishing an equilibrium mixture of SQ anomers. A pertinent example is that of red blood cells in which it is known that glucose importers exhibit a preference for  $\alpha$ -glucose [45], and a potential role for glucomutarotase in a permease system involved in re-establishing this equilibrium has been advanced [34]. For cells grown on SQ glycosides, SQ is released by action of SQases; the immediate product is  $\alpha$ -SQ, and SQMs may be required to enhance conversion of  $\alpha$ -SQ to  $\beta$ -SQ, to match the preference of a downstream enzyme, or to act in the reverse direction to 'rescue'  $\beta$ -SQ that accumulates as a result of the spontaneous

mutarotation of  $\alpha$ -SQ. Downstream enzymes include SQ isomerases and SQ dehydrogenases. As yet the anomeric preference, if any, of these enzymes is unknown. The NMR EXSY experiments used herein to probe SQM activity, which build on pioneering work reaching back many decades, could be of use for studying the substrate preferences of the downstream enzymes, and in so doing could help provide the biological context for the SQMs.

## Abbreviations

DHAP, dihydroxyacetone phosphate; DHPS, 2,3-dihydroxypropanesulfonate; Gal, galactose; Glc, glucose; GlcA, glucuronic acid; Man, mannose; ED, Entner-Doudoroff; EMP, Embden-Meyerof-Parnas; EXSY, exchange spectroscopy; NMR, nuclear magnetic resonance; PNPSQ, 4-nitrophenyl  $\alpha$ -D-sulfoquinovoside; SL, sulfolactate; SQ, sulfoquinovose; SQase, sulfoquinovosidase; SQM, sulfoquinovose mutarotase.

## Author Contribution

S.J.W., S.Y. and E.D.G-B. conceived and coordinated the study. J.P.L. and J.Y. performed the cloning and protein expression. P.A. and S.Y. conducted the NMR measurements. P.A. and C.B. conducted the polarimetry measurements. All authors contributed to writing of the manuscript and approved the final version of the manuscript.

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## Competing Interests

The Authors declare that there are no competing interests associated with the manuscript.

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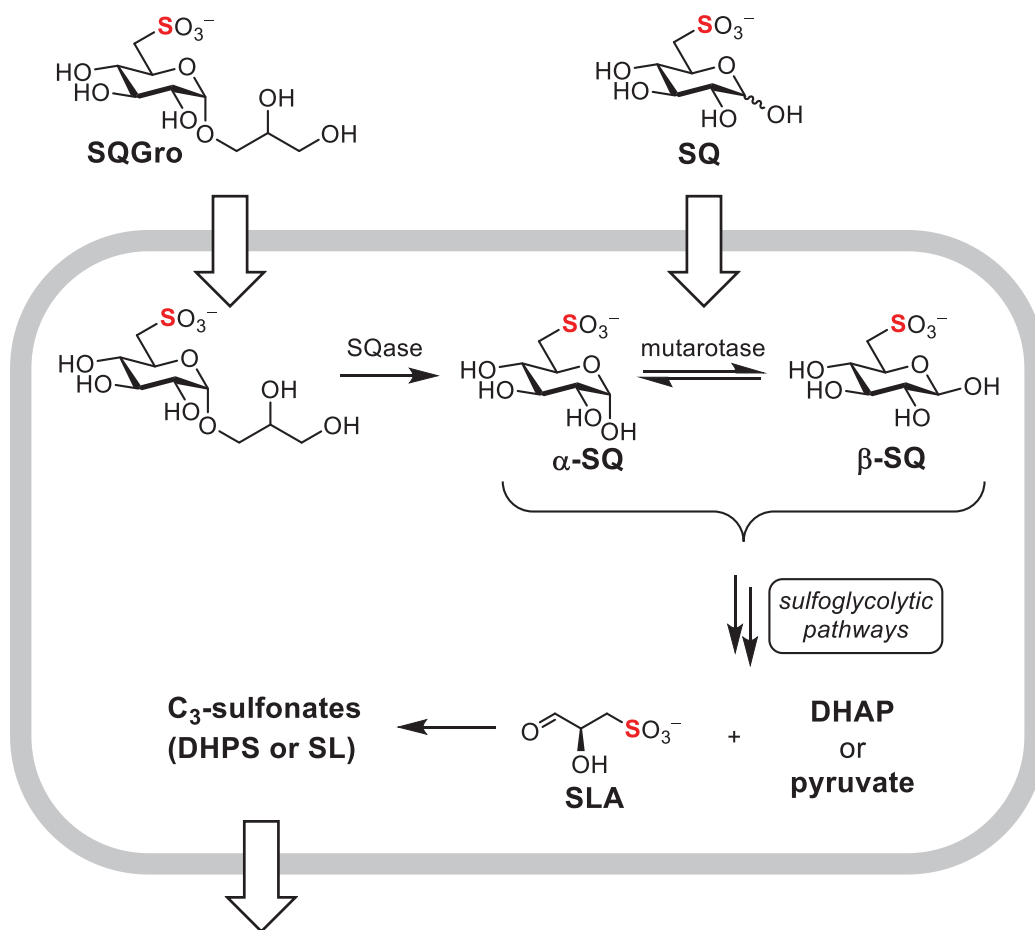
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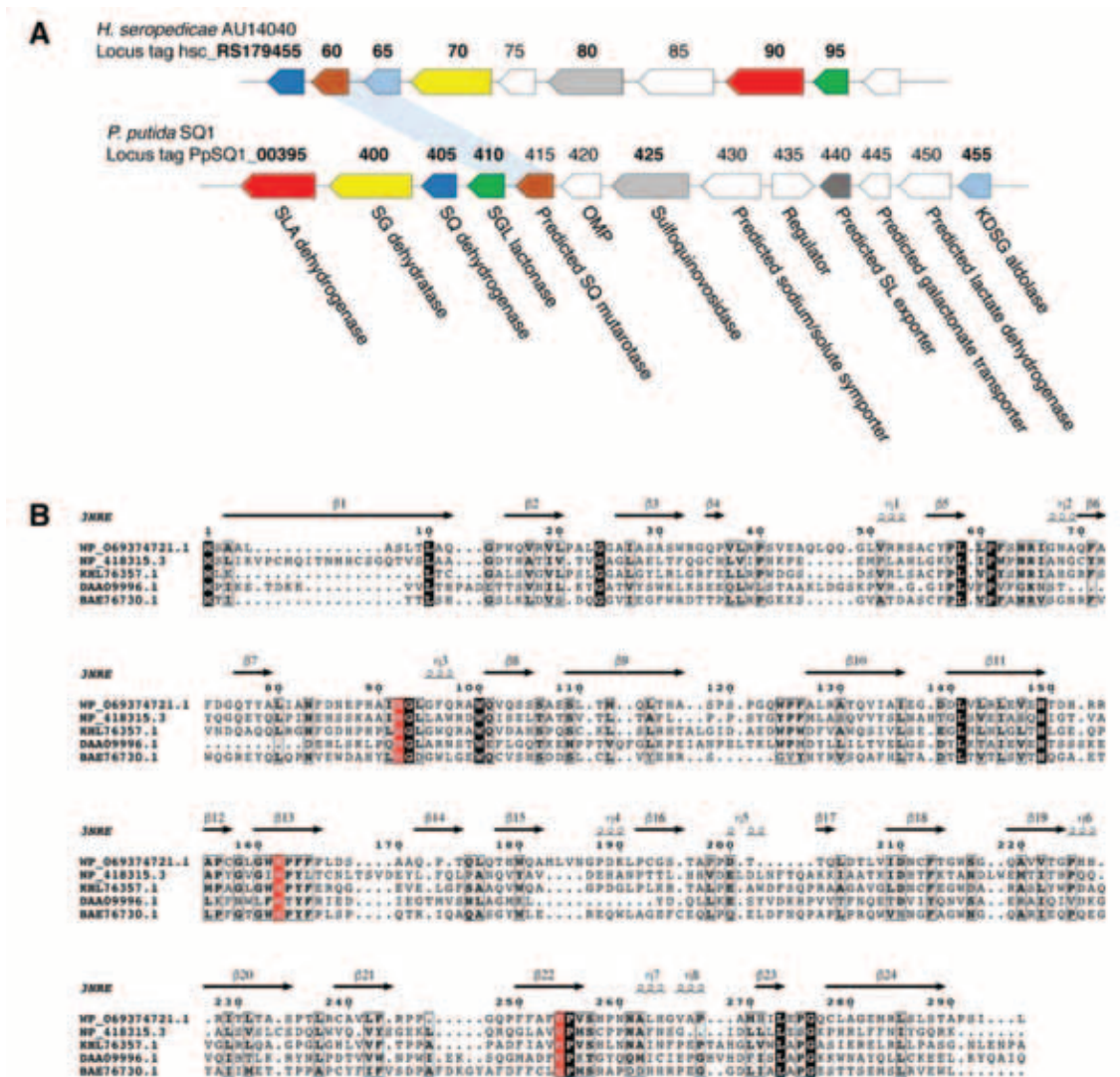


587

588 **Figure 1. Summary of sulfoglycolysis.**

589 Importation of SQGro and cleavage by SQase, or direct importation of SQ, provides an  
 590 intracellular pool of SQ anomers that can be interconverted by SQ mutarotase. SQ is  
 591 metabolized by sulfo-ED or sulfo-EMP pathways to sulfolactaldehyde (SLA), and then to the  
 592 C<sub>3</sub>-sulfonates dihydroxypropanesulfonate (DHPS) or sulfolactate (SL), prior to export.

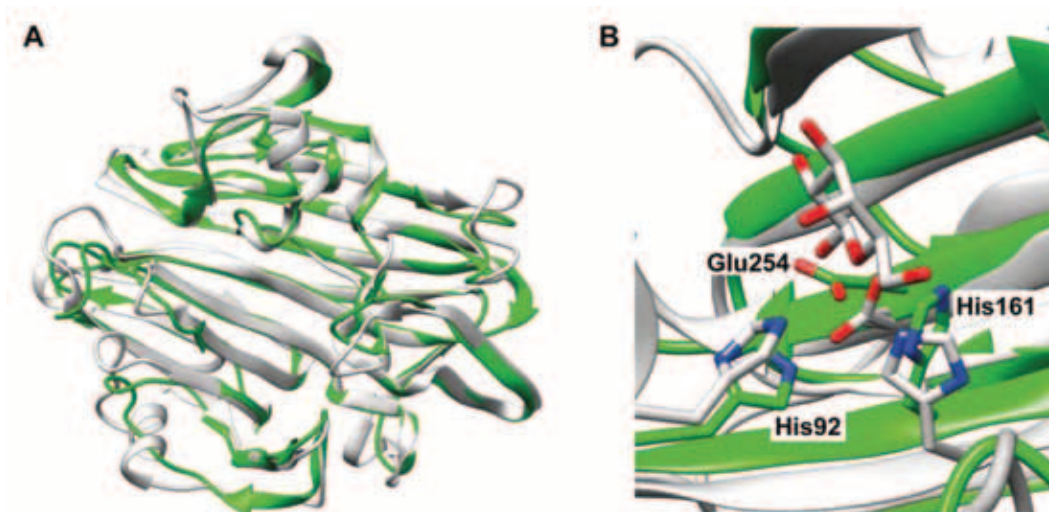
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**Figure 2. *Herbaspirillum seropedicae* contains a sulfo-ED operon and a putative SQ mutarotase.**

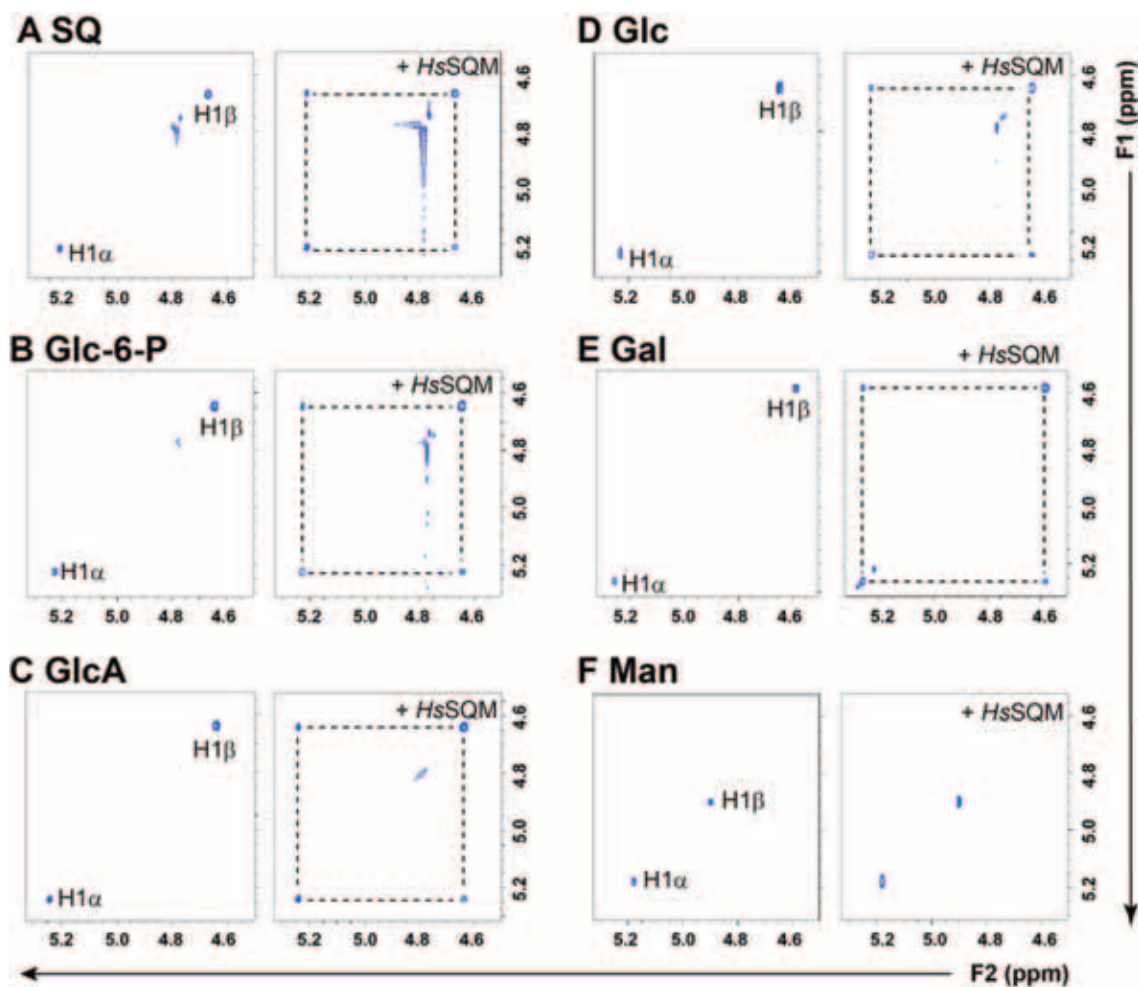
A) Operon structure of *P. putida* SQ1 and *H. seropedicae* strain AU14040. Bold indicates genes for which enzymatic activity has been biochemically determined in at least one organism.

B) Alignment of various putative mutarotases with secondary structural elements. WP\_069374721.1, HsQM from *H. seropedicae*; NP\_418315.3, YihQ from *E. coli*; KHL76357.1, PpSQ1\_00415 from *Pseudomonas putida* SQ1; DAA09996.1, YMR099C hexose-1-phosphate mutarotase from *S. cerevisiae*; BAE76730.1, YphB aldose-1-epimerase from *E. coli* (BAE76730.1). The secondary structural elements are annotated from the structure of *E. coli* YphB (PDB 3nre).



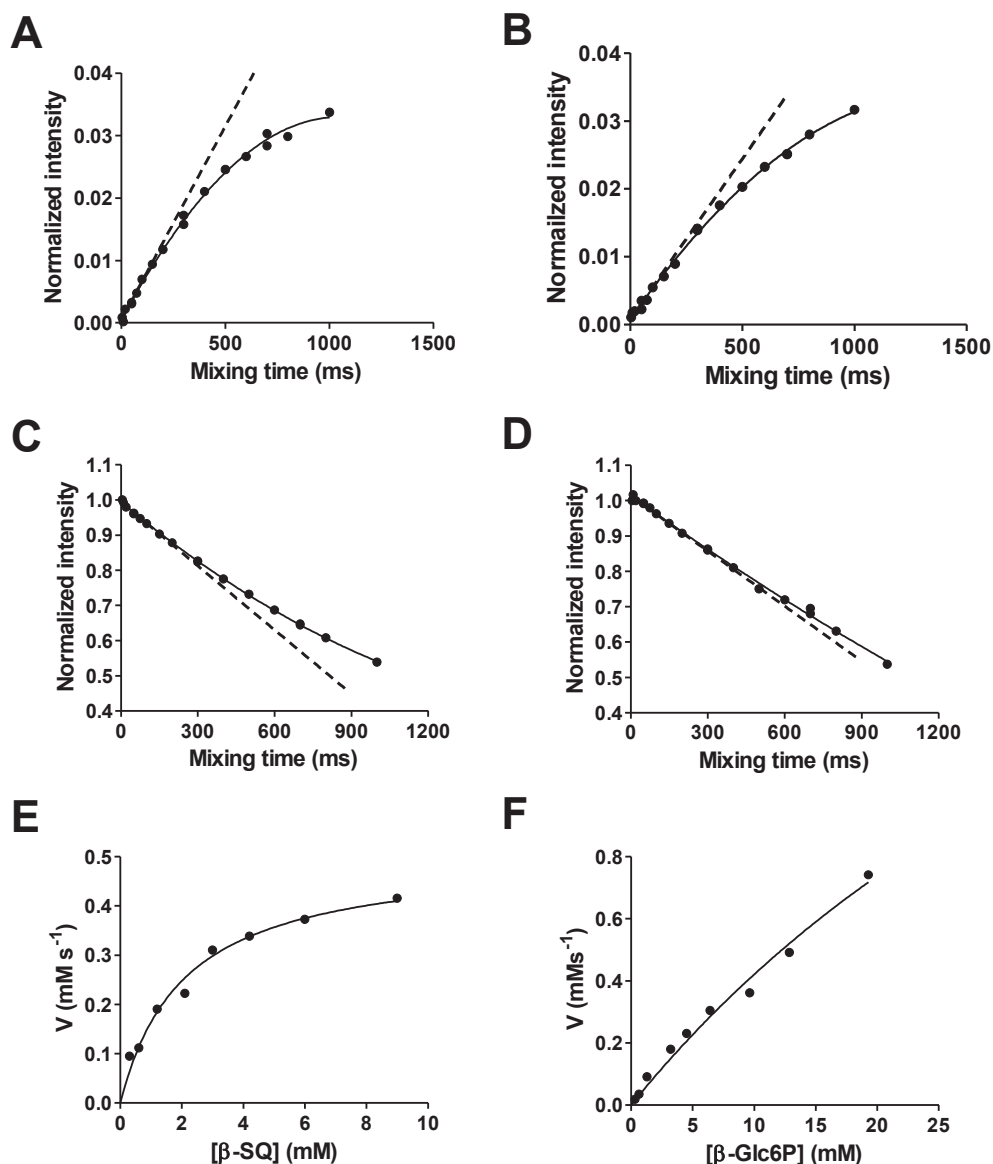
**Figure 3. Homology model of *HsSQM*.**

A) Overlay of *HsSQM* homology model (green) with a putative mutarotase from *Clostridium acetobutylicum* (PDB 3OS7; grey). B) The active sites of the *HsSQM* homology model (green) overlaid with the galactose mutarotase domain of gal10 from *S. cerevisiae* with D-galactose bound (grey, PDB 1Z45). Residue numbers are for *HsSQM*.



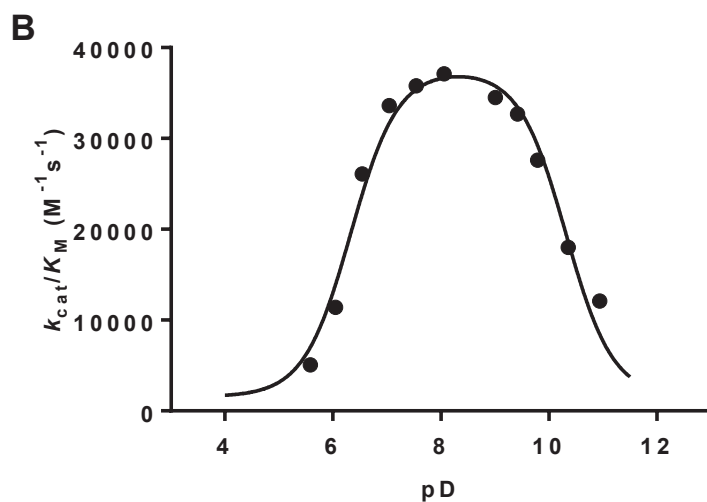
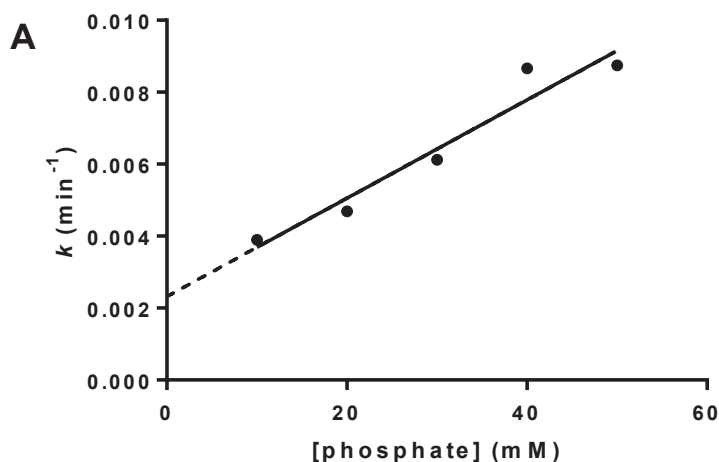
**Figure 4. Excerpt showing anomeric regions of 2D  $^1\text{H}$ - $^1\text{H}$  EXSY plots of various hexoses alone and with *H. seropediacae* mutarotase.**

A) sulfoquinovose (SQ), B) D-glucose-6-phosphate (Glc-6-P), C) D-glucuronic acid (GlcA), D) D-glucose (Glc), E) D-galactose (Gal), F) D-mannose (Man). Hexoses are at 5 mM, 1.51  $\mu\text{M}$  HsSQM in 50 mM sodium phosphate, 150 mM NaCl (pD 7.5).



**Figure 5. Kinetic analysis of *HsSQM* by inversion-recovery 1D  $^1\text{H}$  EXSY.**

Inversion recovery curves for 10 mM SQ or Glc-6-P corresponding to (A)  $\alpha$ -SQ at 4.00 mM and (B)  $\alpha$ -Glc-6-P at 3.57 mM. Inversion decay curves corresponding to (C)  $\beta$ -SQ at 6.00 mM and (D)  $\beta$ -Glc-6-P at 6.43 mM. Michaelis-Menten plots for (E)  $\beta$ -SQ and (F)  $\beta$ -Glc-6-P. Dashed lines indicate tangents to fitted curve at  $t = 0$ .



**Figure 6. Spontaneous mutarotation of  $\alpha$ -SQ and pD dependence of *HsSQM*.**

A) Plot of rates of spontaneous mutarotation of  $\alpha$ -SQ versus phosphate buffer concentration at pseudo-constant ionic strength. Buffers consisted of 10-50 mM sodium phosphate, 2 M NaCl in D<sub>2</sub>O (pD 7.5). For comparison,  $k = 0.0073 \text{ min}^{-1}$  in 50 mM sodium phosphate, 150 mM NaCl in D<sub>2</sub>O (pD 7.5). B) pD dependence of *HsSQM* activity for mutarotation of glucose-6-phosphate. Data was fit to a bell-shaped curve leading to estimated  $pK_a$  values of  $5.9 \pm 0.1$  and  $9.9 \pm 0.1$ .



# Discovery and characterization of a sulfoquinovose mutarotase using kinetic analysis at equilibrium by exchange spectroscopy

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## HsSQM plasmid and sequence data

plasmid	resistance	details
pET29-HsSQM	Kan	C-terminal His <sub>6</sub> -tagged <i>Herbaspirillum seropedicae</i> strain AU14040 sulfoquinovose mutarotase (WP_069374721.1)

### pET29-HsSQM

#### CDS:

ATGTCGGCAGCTCTTGCGTCCCTGACACTTGCCCAAGGACCGTGGCAAGTGCGCGTGCTGCCTGCACTGGGCGGTGCGATTGC  
AAGCGCCAGCTGGCGTGGTCAGCCGGTGCTGCGCCCGAGCGTCGAGGCACAGCTGCAGCAGGGCCTGGTTAGACGCTCTGCGT  
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ACTGA

#### Translation:

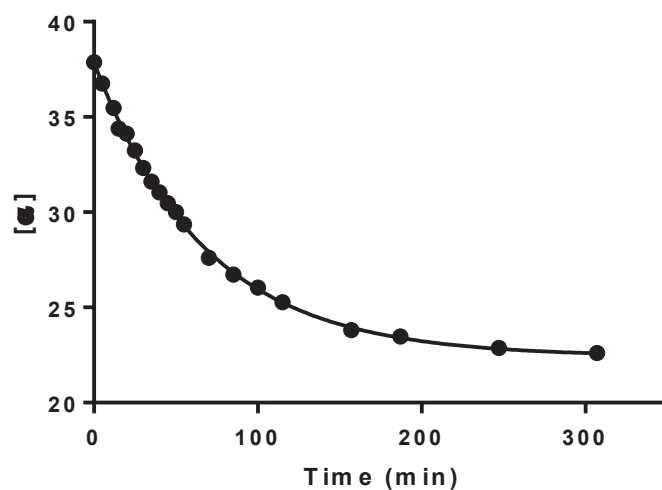
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VLFRPPGQPPFAFEPVSHPNNALHGVPAMHILEPGQCLAGEMRLSLSTAPSILAAALEH  
HHHHH\*

**Table S1. Pairwise sequence comparisons of HsSQM with various hexose mutarotases.\***

<i>H. seropediaceae</i>	<i>P. putida</i>	<i>E. coli</i>	<i>L. lactis</i>	<i>S. cerevisiae</i>	
<b>HsSQM</b> WP_069374721.1	<b>PpSQ_00415</b> KHL76357.1	<b>YihQ</b> NP_418315.3	<b>GalM</b> WP_039116219.1	<b>YMR099C</b> DAA09996.1	<b>% identity</b> <b>(%similarity)</b>
	35 (49)	19 (38)	13 (26)	13 (28)	<b>HsSQM</b>
		19 (37)	12 (24)	15 (29)	<b>PpSQ_00415</b>
			12 (29)	11 (23)	<b>YihQ</b>
				14 (23)	<b>GalM</b>
					<b>YMR099C</b>

\* Calculated using the “Sequence Manipulation Suite” and the following groupings of similar amino acids: GAVLI, FYW, CM, ST, KRH, DENQ, P.

See: [http://www.bioinformatics.org/sms2/ident\\_sim.html](http://www.bioinformatics.org/sms2/ident_sim.html), Stothard, P. *Biotechniques*, 2000, 28, 1102–1104.

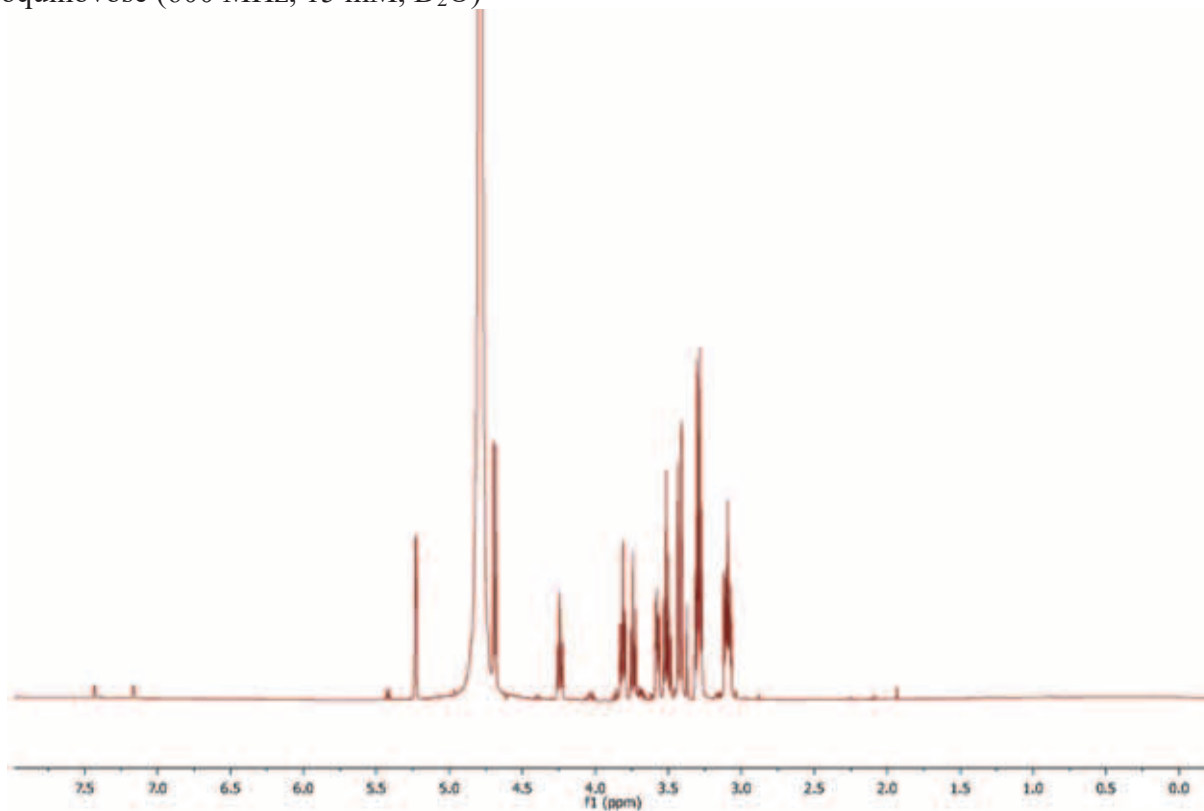


**Figure S1. Mutarotation of SQ monitored by polarimetry.**

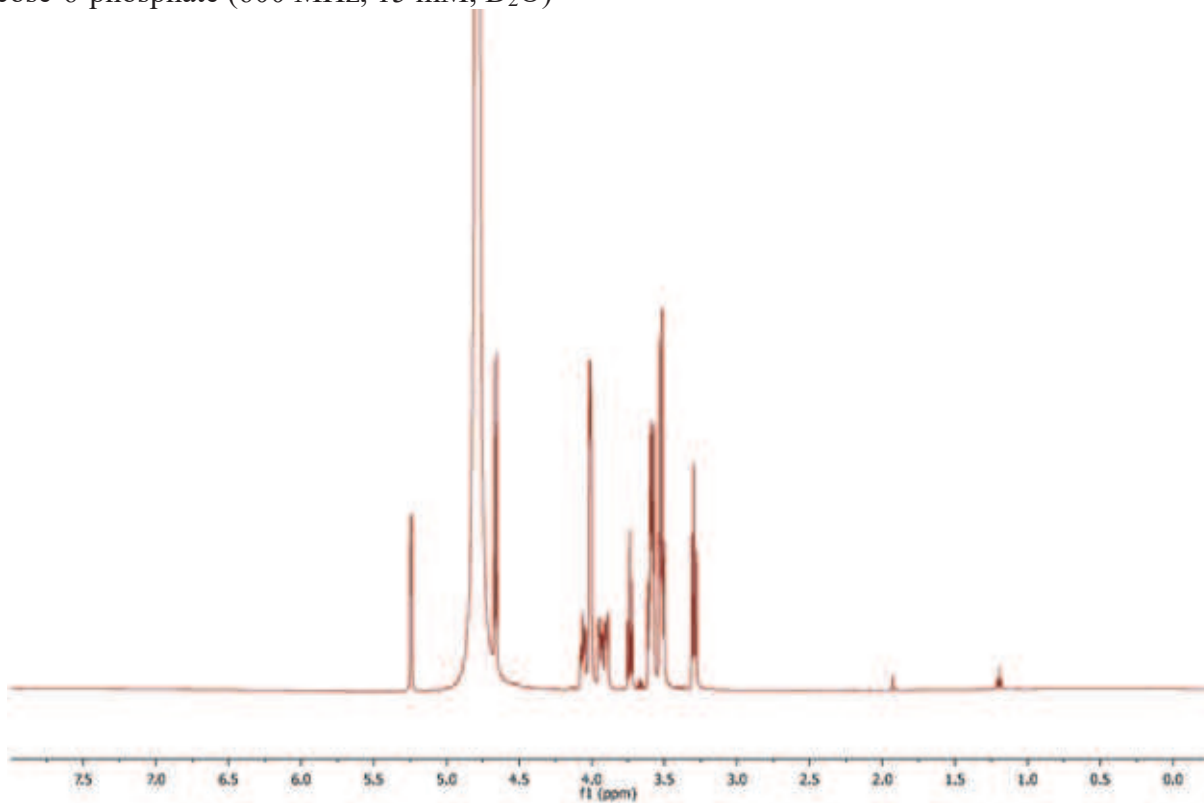
4-Nitrophenyl  $\alpha$ -D-sulfoquinovoside in 50 mM sodium phosphate, 150 mM NaCl in H<sub>2</sub>O (pH 7.1) was treated with sulfoquinovosidase, and after 5 min, the specific rotation was monitored as a function of time. pH 7.1, 26 $\pm$ 1 °C.

# 1D <sup>1</sup>H NMR spectra

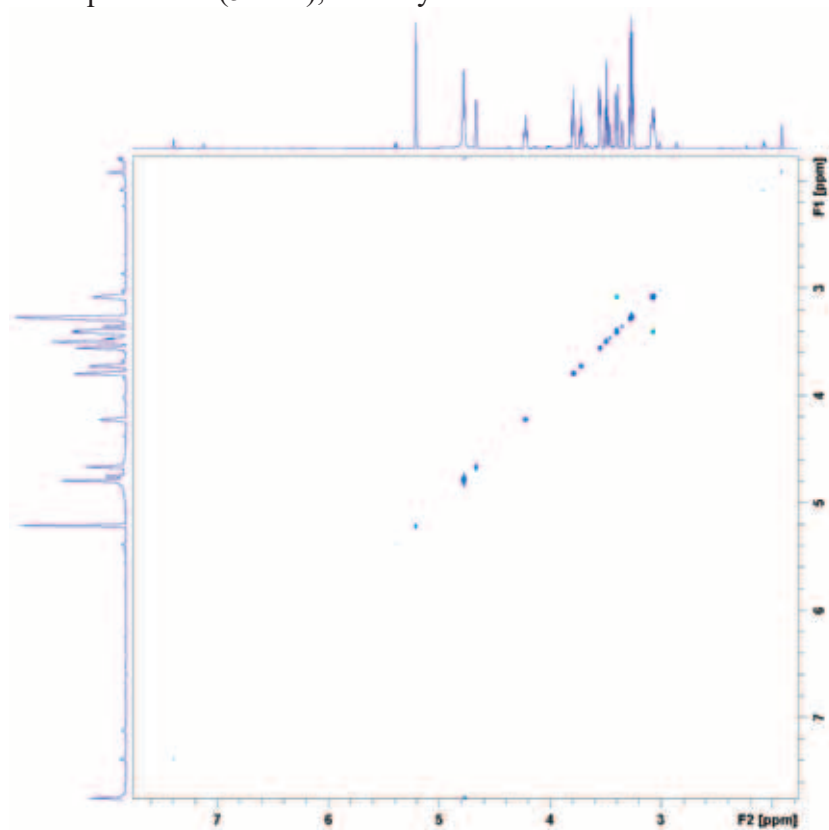
Sulfoquinovose (600 MHz, 15 mM, D<sub>2</sub>O)



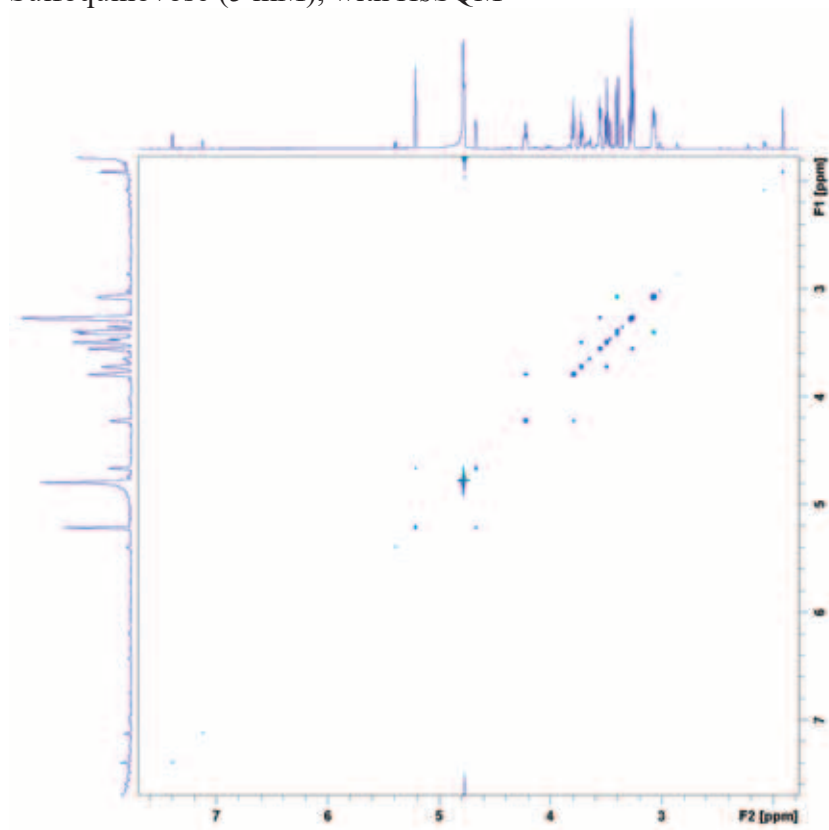
Glucose-6-phosphate (600 MHz, 15 mM, D<sub>2</sub>O)



**2D  $^1\text{H}$ - $^1\text{H}$  NOESY spectra with 1D spectrum projected onto F1 and F2 axes**  
Sulfoquinovose (5 mM), no enzyme

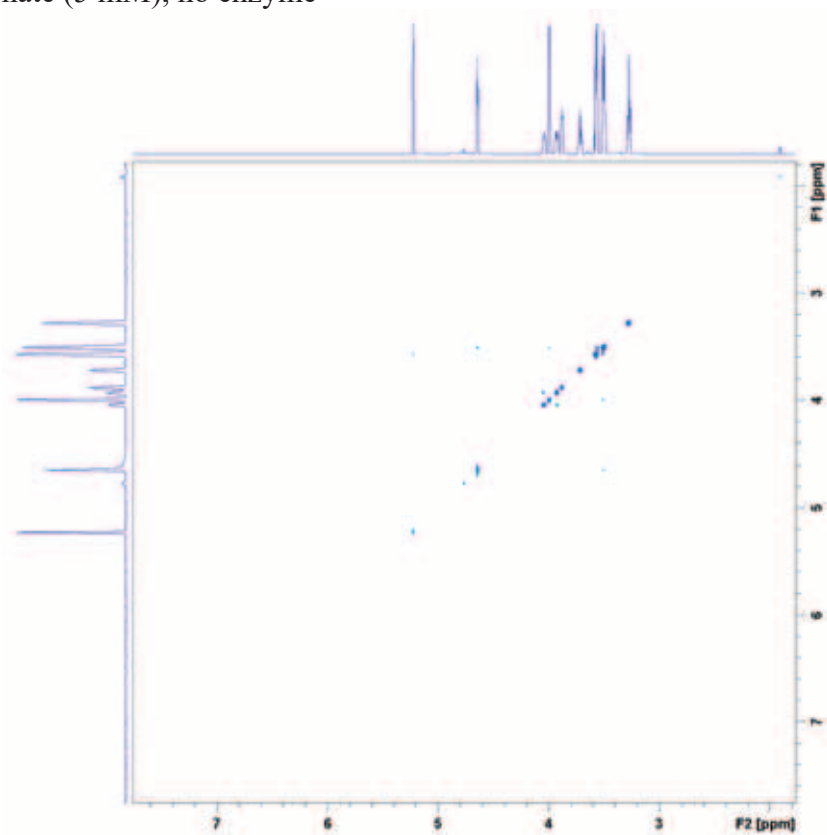


Sulfoquinovose (5 mM), with *HsSQM*

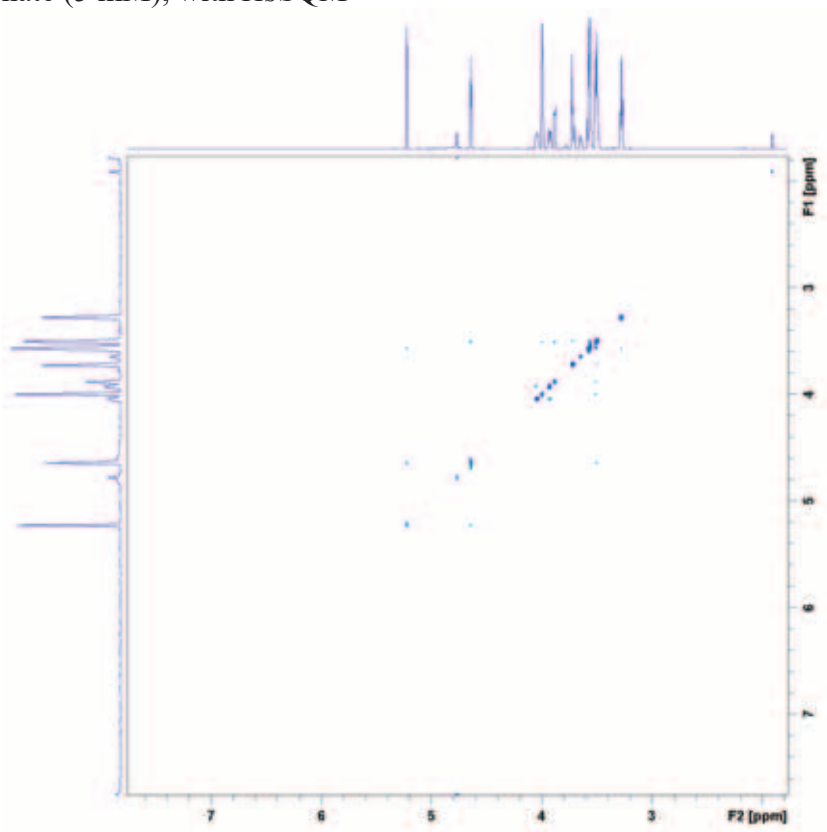




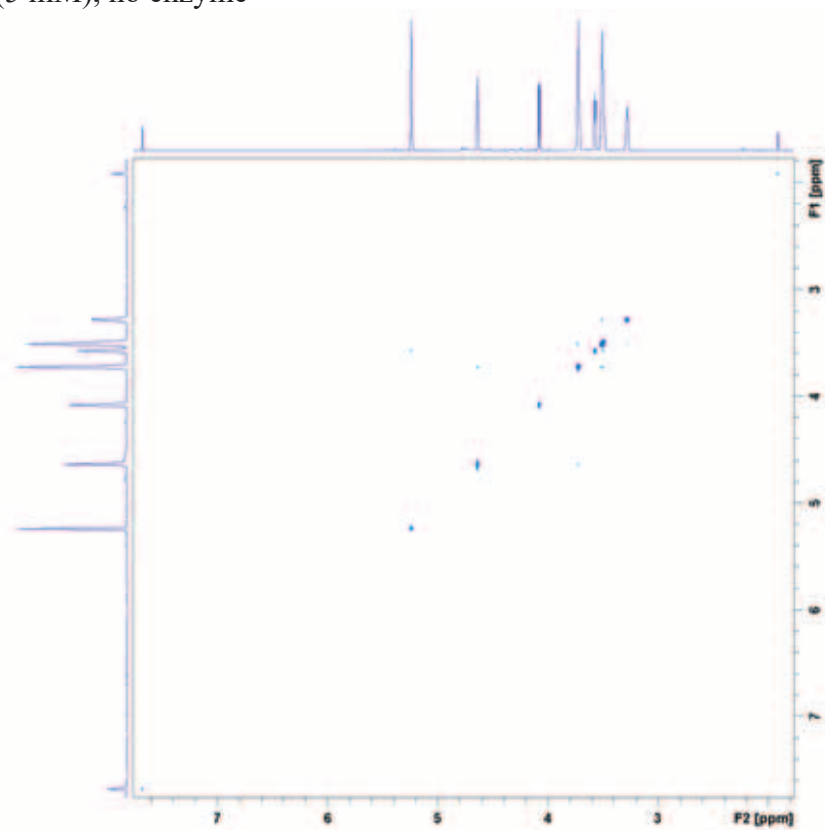
D-Glucose-6-phosphate (5 mM), no enzyme



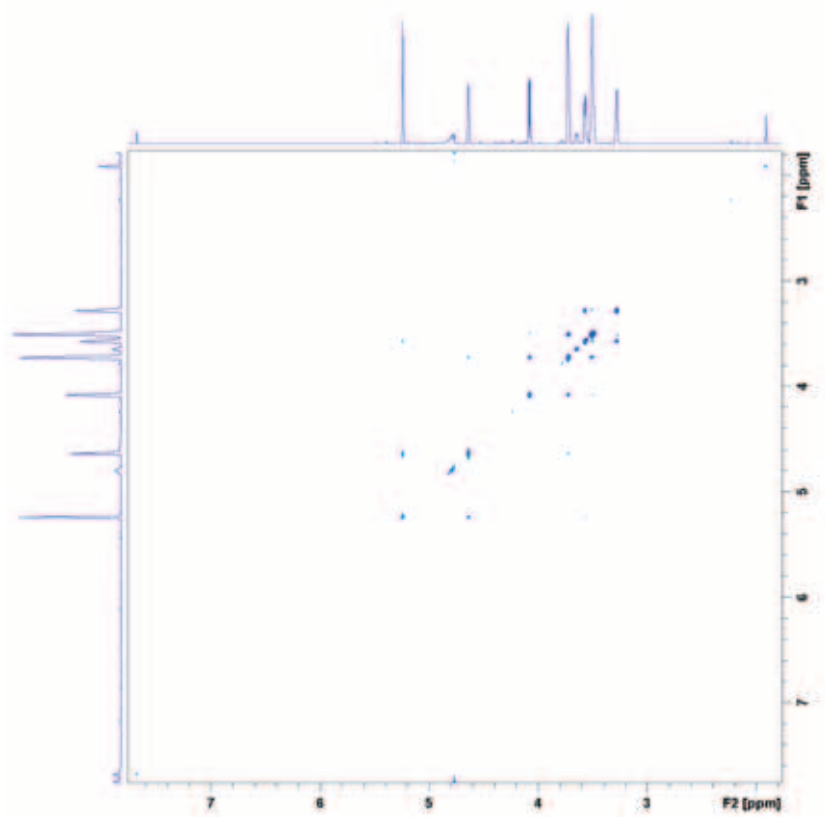
D-Glucose-6-phosphate (5 mM), with *HsSQM*



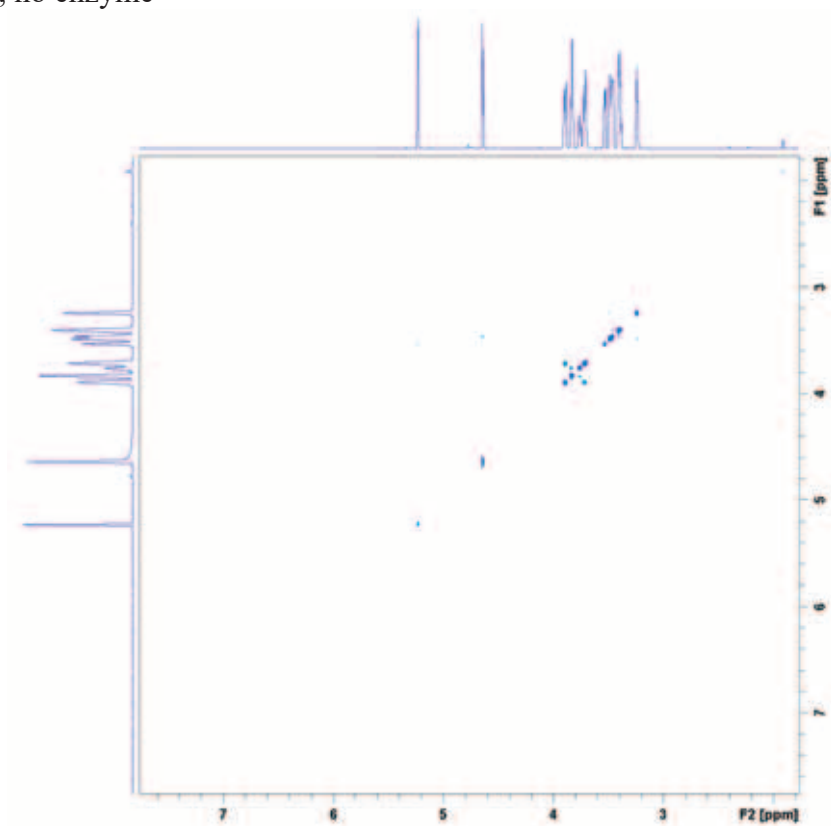
D-Glucuronic acid (5 mM), no enzyme



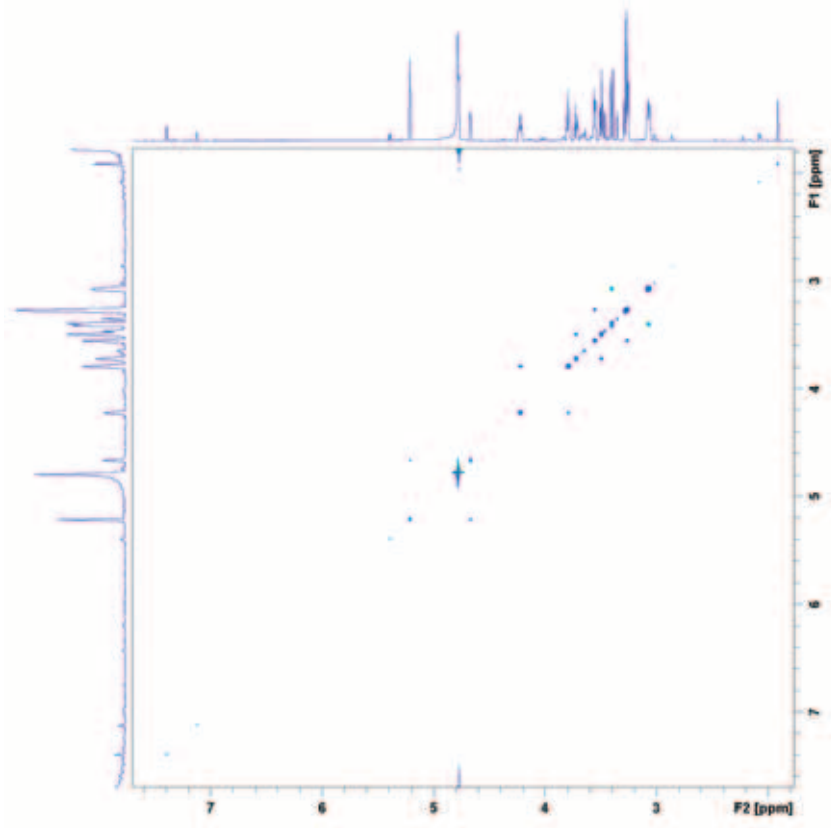
D-Glucuronic acid (5 mM), with *HsSQM*



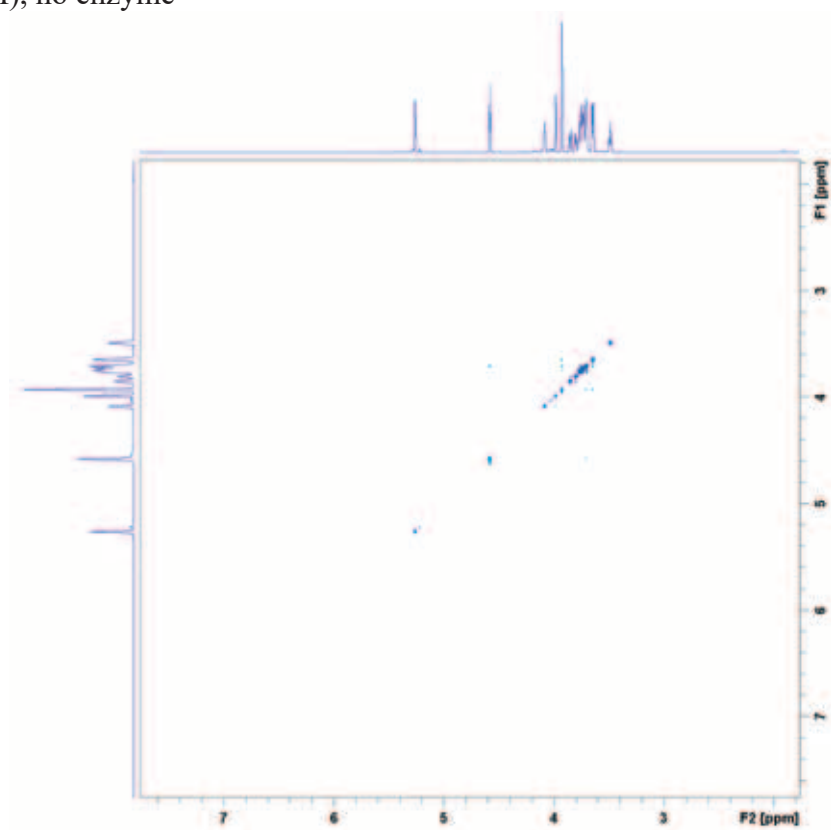
D-Glucose (5 mM), no enzyme



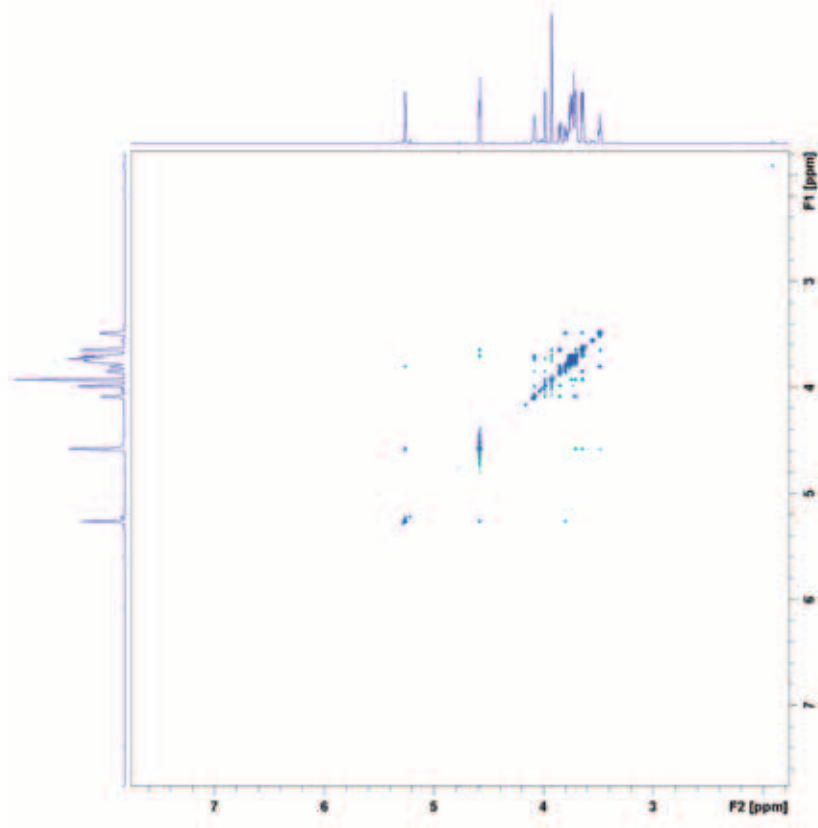
D-Glucose (5 mM), with *HsSQM*



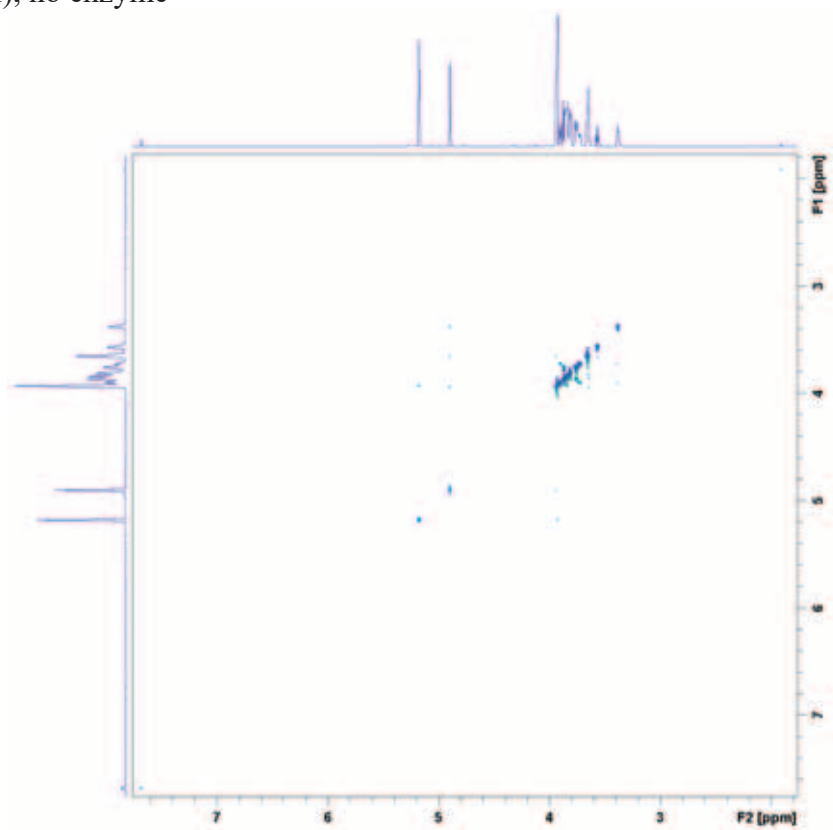
D-Galactose (5 mM), no enzyme



D-Galactose (5 mM), with *HsSQM*



D-Mannose (5 mM), no enzyme



D-Mannose (5 mM), with *HsSQM*

